

U.S.S.N. 10/027,248  
Attorney Docket No.: BAI-007CPACN

Examiner: S. Tran  
Group Art Unit: 1615

- 032
8. [Amended] A pharmaceutical tablet composition comprising:
- (a) an effective amount of ibuprofen wherein the weight of the ibuprofen is provided in a range, of the total weight of the tablet composition, of up to about 50%;
  - (b) an effective amount of hydrocodone;
  - (c) colloidal silicon dioxide provided in a range, by total weight of the tablet composition, of about 1.5% to about 2%;
  - (d) microcrystalline cellulose provided in a range, of the total weight of the tablet composition, of about 15% to about 25%;
  - (e) a disintegrant selected from the group consisting of croscarmellose sodium and crospovidone wherein the weight of the disintegrant is provided in a range, of the total weight of the tablet composition, of about 6 to about 8%;
  - (f) a binder consisting of an alkylhydroxy methylcellulose wherein the weight of the binder is provided in a range, of the total weight of the tablet composition, of about 3 % to about 4 %;
  - (g) corn starch wherein the weight of the corn starch is provided in a range, of the total weight of the tablet composition, of about 11 to about 17 %; and
  - (h) a lubricant wherein the weight of the lubricant is provided in an amount less than 1% by weight of the total weight of the tablet.